

Helobdella blinni sp. n. (Hirudinida, Glossiphoniidae) a new species inhabiting Montezuma Well, Arizona, USA

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Academic editor: S. James | Received 30 June 2016 | Accepted 1 March 2017 | Published 15 March 2017

http://zoobank.org/963C2520-3C29-419B-A5D1-0518E9713DD5

Citation: Beresic-Perrins RK, Govedich FR, Banister K, Bain BA, Rose D, Shuster SM (2017) *Helobdella blinni* sp. n. (Hirudinida, Glossiphoniidae) a new species inhabiting Montezuma Well, Arizona, USA. ZooKeys 661: 137–155. https://doi.org/10.3897/zookeys.661.9728

Abstract

A new leech species *Helobdella blinni* **sp. n.**, is described from Montezuma Well, an isolated travertine spring mound located in central Arizona, USA. In its native habitat, *H. blinni* had been previously identified as *Helobdella stagnalis* (Linnaeus, 1758), which was later reclassified to *Helobdella modesta* (Verrill, 1872). Similar to the European *H. stagnalis* and North American *H. modesta*, *H. blinni* has six pairs of testisacs, five pairs of smooth crop caecae, one lobed pair of posteriorly-directed crop caecae, one pair of eyes, a nuchal scute, and diffuse salivary glands. However, the pigmentation of this new species ranges from light to dark brown, unlike *H. modesta* which tends to be light grey in color. Also, *H. modesta* produces a clutch of 12—35 pink eggs, whereas *H. blinni* produces smaller clutches of white eggs (7–14, 0.5 ± 0.15 mm, N = 7) and consequently broods fewer young (1–14, 7 ± 3.3 mm, N = 97). *Helobdella blinni* are also able to breed year-round due to the constant warm water conditions in Montezuma Well. Their breeding season is not restricted by seasonal temperatures. These species are morphologically similar, however, comparing the COI mtDNA sequences of *H. blinni* with sequences from nearby populations of *H. modesta* and other *Helobdella* species from GenBank indicate that *H. blinni* is genetically distinct from these other *Helobdella* populations.

Keywords

Leech, Glossiphoniidae, Helobdella blinni sp. n., new species, Montezuma Well

Introduction

Montezuma Well is a collapsed travertine spring mound located 72 km south of Flagstaff in the Verde Valley of Northern Arizona (34.6491°N,111.7522°W (DD)) (Fig. 1A). The age of Montezuma Well is estimated to be ~11,000 years (Wagner and Blinn 1987). This location is thermally constant year-round (19–24°C) and is continuously replenished by two vents located at the well bottom. Montezuma Well is 0.76 ha in area and approximately 20 m deep. Most of the shoreline drops off immediately into open water, except at the northeast corner where water drains through a shallow region called the "swallet" and empties into Wet Beaver Creek which is located east of Montezuma Well (Fig. 1A–B). The water within Montezuma Well has unique water chemistry, containing high levels of arsenic (>100μg/L) and dissolved CO₂ (>300mg/L) (Cole and Barry 1973).

Four leech species are known to inhabit Montezuma Well, including an endemic pelagic predator (Govedich et al. 1998), the erpobdellid *Motobdella montezuma* (Davies et al. 1985), and three other glossiphoniid species currently identified as *Helobdella papillata* (Moore, 1952), *H. elongata* (Castle, 1900), and a species currently thought to be *H. stagnalis* (Linnaeus, 1758), all of which inhabit the swallet (Fig. 1B). These Montezuma Well leech populations are thought to have been isolated from other leech populations for as long as 11,000 years (Wagner and Blinn 1987).

In support of this hypothesis, Beresic-Perrins's (2010) description of brood size, parental behavior, and life history of the Montezuma Well population of *H. stagnalis* suggests that this leech is distinct from other known populations of *H. stagnalis*, a species originally described from Europe and which had until very recently been considered to be a widespread cosmopolitan leech species, inhabiting both Europe and North America. Siddall et al. (2005) addressed this problem by resurrecting the original species description for the North American leech, *Helobdella modesta* (Verrill, 1872) which had long been considered to be a synonym of the European *H. stagnalis* (Moore 1898). The molecular analysis by Moser et al. (2011) provided confirmation for the resurrection of *H. modesta* by Siddall et al. (2005). Even though the two species are morphologically indistinguishable (Verrill 1872, Moore 1898, Moore 1952), they differ genetically. Henceforth, we will refer to the North American *H. stagnalis* as *H. modesta*.

Here, we compare key traits, both morphological and molecular, among members of the Montezuma Well *Helobdella* sp. population, several other nearby populations of *H. modesta*, and several other *Helobdella* species. Our molecular analysis includes the cytochrome *c* oxidase subunit I (COI) mitochondrial gene region to test the hypothesis that the Montezuma Well population of *H. modesta* is a distinct species and warrants a new species description. This region is known to be sufficiently variable to reveal interspecific differences and unlikely to suggest differences due to elevated mutation rates (Apakupakul et al. 1999).

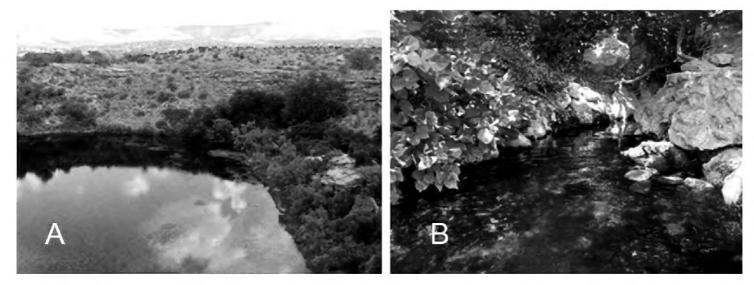


Figure 1. Location of *Helobdella blinni* sp. n. **A** The northeast side of Montezuma Well; and **B** The swallet where the leeches were collected.

Materials and methods

Sampling

A total of 34 individuals of *Helobdella* sp. inhabiting Montezuma Well were collected from the underside of rocks in the swallet: five specimens were collected in June 2011 for molecular analysis and 29 were collected in June 2012 to assess morphological characteristics. For the molecular analysis, the leeches were preserved in 95% ethanol and others, for museum collections, were fixed with buffered formalin overnight and preserved in 70% ethanol. Additionally, a total of 10 specimens of *H. c.f. modesta* from Rio de Flag ponds near the Rio de Flag Waste Water Facility outflow in Flagstaff, Arizona (35.18418°N, 111.63294°W (DD)) and Oak Creek, AZ near the Cave Springs campground (34.9961°N, 111.7394°W (DD)) were collected for molecular analyses. These specimens were also fixed in 95% ethanol.

Morphological examination

We documented number of eyes and their placement, color pattern, presence of papillae, number of and structure of gastric caecae, body size, presence of nuchal scute, gonopore placement, egg size and number, and number of offspring using a Nikon binocular dissecting microscope. We then deposited the examined materials in the Invertebrate Zoology collection at the Smithsonian Institution, National Museum of Natural History (USNM).

Molecular analysis

Whole DNA was extracted from the caudal suckers of the individual leeches using a Qiagen DNeasy Blood & Tissue Kit (Cat. No. 69504), with each sample incubated

Table 1. Helobdella and outgroup taxa used for our molecular analysis.

Taxon	Locality	Reference
Cystobranchus salmositicus	Outgroup	Williams and Burreson 2006
Ozobranchus margoi	Outgroup	Siddall and Burreson 1998
Gonimosobdella klemmi	Outgroup	Williams and Burreson 2005
Myzobdella lugubris	Outgroup	Siddall and Burreson 1998
Helobdella atli	French Guiana	Oceguera-Figueroa et al. 2010
Helobdella atli	Uruguay	Oceguera-Figueroa et al. 2010
Helobdella atli	Mexico	Oceguera-Figueroa et al. 2010
Helobdella blinni sp. n.	Montezuma Well, AZ, USA	This study
Helobdella blinni sp. n.	Montezuma Well, AZ, USA	This study
Helobdella blinni sp. n.	Montezuma Well, AZ, USA	This study
Helobdella bolivianita	Bolivia	Siddall and Borda 2002
Helobdella bowermani	Oregon, USA	Moser et al. 2013
Helobdella bowermani	Oregon, USA	Moser et al. 2013
Helobdella bowermani	Oregon, USA	Moser et al. 2013
Helobdella californica	California, USA	Kutschera 2011
Helobdella "elongata"	Mexico	Oceguera-Figueroa et al. 2010
Helobdella europaea	Taiwan	Lai et al. 2009
Helobdella europaea	Taiwan	Lai et al. 2009
Helobdella europaea	Taiwan	Lai et al. 2009
Helobdella europaea	Taiwan	Lai et al. 2009
Helobdella europaea	South Africa	Siddall and Budinoff 2005
Helobdella lineata	Michigan, USA	Siddall and Borda 2002
Helobdella fusca	Michigan, USA	Siddall and Borda 2002
Helobdella melananus	Taiwan	Lai et al. 2009
Helobdella melananus	Taiwan	Lai et al. 2009
Helobdella melananus	Taiwan	Lai et al. 2009
Helobdella michaelseni	Chile	Siddall and Borda 2002
Helobdella modesta	Columbus, Ohio, USA	Siddall and Borda 2002
Helobdella modesta	Washington, USA	Oceguera-Figueroa et al. 2010
Helobdella modesta	Washington, USA	Oceguera-Figueroa et al. 2010
Helobdella c.f. modesta	Rio de Flag, Flagstaff, AZ, USA	This study
Helobdella c.f. modesta	Rio de Flag, Flagstaff, AZ, USA	This study
Helobdella c.f. modesta	Oak Creek, AZ, USA	This study
Helobdella c.f. modesta	Oak Creek, AZ, USA	This study
Helobdella nununununojensis	Bolivia	Siddall and Borda 2002
Helobdella nununununojensis	Bolivia	Siddall and Borda 2002
Helobdella octatestisaca	Taiwan	Lai et al. 2009
Helobdella octatestisaca	Taiwan	Lai et al. 2009
Helobdella octatestisaca	Taiwan	Lai et al. 2009
Helobdella octatestisaca	Taiwan	Lai et al. 2009
Helobdella octatestisaca	Taiwan	Lai et al. 2009
Helobdella octatestisaca	Taiwan	Lai et al. 2009
Helobdella octatestisaca	Taiwan	Lai et al. 2009
Helobdella octatestisaca	South Africa	
rreiovaeua ociaiestisaca	Souul Airica	Oceguera-Figueroa et al. 2010

Taxon	Locality	Reference
Helobdella octatestisaca	Mexico	Oceguera-Figueroa et al. 2010
Helobdella octatestisaca	Mexico	Oceguera-Figueroa et al. 2010
Helobdella octatestisaca	Mexico	Oceguera-Figueroa et al. 2010
Helobdella octatestisaca	Mexico	Oceguera-Figueroa et al. 2010
Helobdella octatestisaca	Mexico	Oceguera-Figueroa et al. 2010
Helobdella papillata	Michigan, USA	Siddall and Borda 2002
Helobdella papillata	Virginia, USA	Siddall and Borda 2002
Helobdella papillornata	Australia	Siddall and Borda 2002
Helobdella paranensis	Uruguay	Siddall and Borda 2002
Helobdella pichipanan	Bolivia	Siddall et al. 2005
Helobdella "robusta" TXAU1	Texas, USA	Bely and Weisblat 2006
Helobdella "robusta"	California, USA	Bely and Weisblat 2006
Helobdella "robusta" CASA 1	California, USA	Bely and Weisblat 2006
Helobdella "robusta" NYTA	New York, USA	Bely and Weisblat 2006
Helobdella simplex	Argentina	Moser et al. 2006
Helobdella simplex	Argentina	Moser et al. 2006
Helobdella simplex	Argentina	Moser et al. 2006
Helobdella socimulcensis	Mexico	Oceguera-Figueroa et al. 2010
Helobdella socimulcensis	Mexico	Oceguera-Figueroa et al. 2010
Helobdella socimulcensis	Mexico	Oceguera-Figueroa et al. 2010
Helobdella socimulcensis	Mexico	Oceguera-Figueroa et al. 2010
Helobdella socimulcensis	Mexico	Oceguera-Figueroa et al. 2010
Helobdella socimulcensis	Mexico	Oceguera-Figueroa et al. 2010
Helobdella socimulcensis	Mexico	Oceguera-Figueroa et al. 2010
Helobdella sp. Xochimilco	Mexico	Oceguera-Figueroa et al. 2010
Helobdella sorojchi	Bolivia	Siddall and Borda 2002
Helobdella sorojchi	Bolivia	Siddall and Borda 2002
Helobdella stagnalis	United Kingdom	Siddall and Borda 2002
Helobdella "stagnalis"	Mexico	Oceguera-Figueroa et al. 2010
Helobdella "stagnalis"	Mexico	Oceguera-Figueroa et al. 2010
Helobdella transversa	Michigan, USA	Siddall and Borda 2002
Helobdella triserialis	Bolivia	Siddall and Borda 2002
Helobdella triserialis	California, USA	Bely and Weisblat 2006
Helobdella virginiae	Mexico	Oceguera-Figueroa et al. 2010

overnight in a water bath set at 54°C. Using Siddall and Borda's (2002) PCR method, the mitochondrial gene region, cytochrome c oxidase subunit I (COI) was amplified. The primers were LCO1490 5'-GGTCAACAAATCATAAAGATATTGG-3' and HCO2198 5'-TAAACTTCAGGGTGACCAAAAAATCA-3' (Folmer et al. 1994). The PCR product was purified through the use of the QIAquick PCR Purification Protocol (Cat. No. 28104), checked for PCR product using gel electrophoresis, and sequenced with an ABI Prism 3730 sequencer (Applied Biosystems). We imported the seven "cleanest" sequences and 71 comparative sequences (Table 1) from previous studies (Siddall and Burreson 1998, Siddall and Borda 2002, Siddall and Budinoff

2005, Siddall et al. 2005, Williams and Burreson 2005, Bely and Weisblat 2006, Williams and Burreson 2006, Lai et al. 2009, Oceguera-Figueroa et al. 2010, Kutschera 2011, Moser et al. 2013) from GenBank (http://www.ncbi.nlm.nih.gov/genbank/) into MEGA7.0.18 (Kumar et al. 2016). We aligned the sequences automatically using MUSCLE (Edgar 2004) and then corrected the alignments by hand. We partitioned the data and performed the substitution model test by codon in Partitionfinder (Lanfear et al. 2012). The best substitution model test was General Time Reversal (GTR) +gamma which we used in our maximum-likelihood (ML) analysis (Lanave et al. 1984, Tavare 1986, Rodriguez et al. 1990). For ML analysis, we used RAxML v. 8 (Stamatakis 2014) and included 1,000 nonparametric bootstrap replicates. We used MrBayes for Bayesian inference analysis with ten million generations with a 25% burn-in and our support was assessed based on clade posterior probabilities (Ronquist and Huelsenbeck 2003). These analyses were conducted through CIPRES (Miller et al. 2010). We used PAUP* 4.0 (Swofford 2003) to construct parsimony phylogenies with 100 random additions. We performed the parsimony analysis twice, treating the deletions in the sequences as a 5th state and then as missing data. We performed an uncorrected p-distance analysis to examine nucleotide differences between sequences with 1,000 replicates in MEGA7.0.18 (Kumar et al. 2016).

Results

Family Glossiphoniidae Vaillant, 1890 Genus *Helobdella* Blanchard, 1896

Helobdella blinni sp. n.

http://zoobank.org/B1B3D234-BC3F-4126-BF25-52DA00BA7EB9 Figs 2, 3, 4

Type materials. Holotype. USNM 1186106 (Table 2).

Additional materials. Paratypes. (14 specimens) (USNM 1186107, 1186108, 1186109, 1186110, 1186111, 1186112, 1186113, 1186114, 1186115, 1186116, 1186117, 1186118, 1186119, 1186120) (Table 2)

Type locality. USA, Arizona: Yavapai County, Montezuma Well (34.6491°N, 111.7522°W (DD)), aquatic system, under rocks, 10 June 2012, R.K. Beresic-Perrins.

Etymology. We have named this new species, *Helobdella blinni* in honor of Dr. Dean W. Blinn for his dedication to natural history research at Montezuma Well. For over 20 years at Northern Arizona University, Dr. Blinn studied a wide range of organisms and their interactions at Montezuma Well including predator-prey interactions between *Motobdella montezuma* and the endemic amphipod, *Hyalella montezuma* Cole & Watkins, 1977.

Description. External morphology. Length of specimens 11 to 22 mm (mean + SE 16.6 + 3.2 N=24) and width 3 to 8 mm (5.7 + 1.1 N=28) (Table 3, Figs 2, 3).

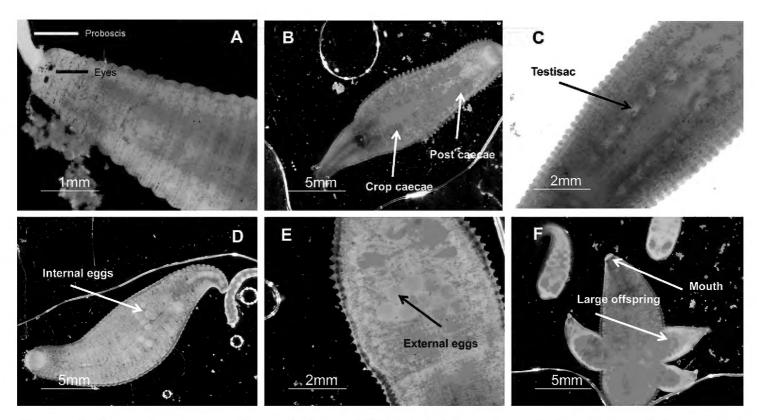


Figure 2. Internal and external morphology of *Helobdella blinni* sp. n. **A** dorsal view of the eyes and extended proboscis **B** crop and post caecae **C** testisacs **D** ventral view of internal eggs which have not been oviposited yet **E** ventral view of white eggs that have been oviposited **F** ventral view of attached and detached offspring.

Table 2. Holotype and paratype collection data and voucher numbers.

Family	Species	Collection data	Voucher #
Glossiphoniidae	Helobdella blinni sp. n.	USA: AZ: Yavapai Co., Montezuma Well 34.6491°N, 111.7522°W (DD), 10.VI.2010, aquatic system, under rocks, RK Beresic-Perrins, Holotype (USNM)	1186106
			1186107
			1186108
			1186109
			1186110
			1186111
		(14 specimens) USA: AZ Yavapai Co.,	1186112
Classinhaniidaa	Ualah dalla hlimai an n	Montezuma Well 34.6491°N, 111.7522°W	1186113
Glossiphoniidae	Helobdella blinni sp. n.	(DD), 10.VI.2010, aquatic system, under	1186114
		rocks, RK Beresic-Perrins, Paratypes (USNM)	1186115
			1186116
			1186117
			1186118
			1186119
			1186120

Individual color ranges from translucent with brown spots to dark brown (Fig. 4). No dorsal papillae; one pair of eyes located at somite II (0.07 + 0.02 mm) diameter, N = 11, distance between eyes 0.1 to 0.03 mm apart (N = 13). A scallop-shaped

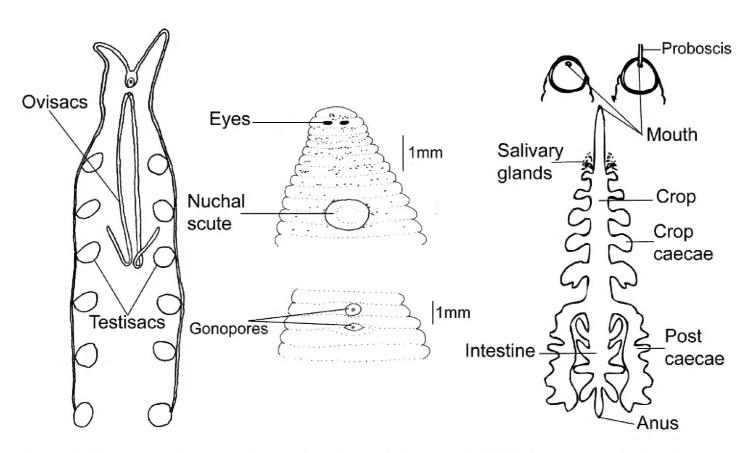


Figure 3. Diagram of the external and internal morphology of *Helobdella blinni* sp. n. (drawn by Rebecca Beresic-Perrins and Fredric Govedich).

Table 3. Morphological measurements of *Helobdella blinni* sp. n.

Trait	Ave	SE	Min	Max	N
body length relaxed (mm)	16.6	3.18	11.3	22.5	24
body width relaxed (mm)	5.7	1.15	3.1	8.0	28
caudal diameter (mm)	1.7	0.3	1.0	2.3	27
egg diameter (mm)	0.5	0.15	0.2	0.7	28
gonopore separation (mm)	0.1	0.08	0.1	0.3	13
nuchal scute length (mm)	0.335	0.05	0.284	0.432	9
nuchal scute width (mm)	0.32	0.04	0.27	0.386	9
proboscis length (mm)	3.5	1.10	2.0	6.2	17
oral sucker diameter (mm)	0.7	0.19	0.4	1.0	15
progeny length (mm)	3.6	1.68	1.6	6.6	18
progeny width (mm)	1.5	0.8	0.7	2.9	18
# eggs	10.0	2.73	7.0	16.0	7
# progeny	7.2	3.35	1.0	14.0	97
eye diameter (mm)	0.1	0.02	0.0	0.1	11
eye distance (mm)	0.1	0.04	0.0	0.2	13

nuchal scute is present on the dorsal side, length 0.293 to 0.432 mm $(0.335 + 0.05 \, \text{N=9})$ and width 0.27 to 0.386 mm $(0.32 + 0.04 \, \text{N=9})$. One annulus separates the female and male gonopores. The caudal sucker diameter averages 1.6 + 0.3 mm (N = 27). The eggs (diameter 0.5 + 0.15 mm, N = 28) are laid on the ventral side of the parent in soft-walled transparent cocoons (7–11 eggs per cocoon, N = 7). The mouth is located subterminally in the oral sucker (Figs 2, 3).

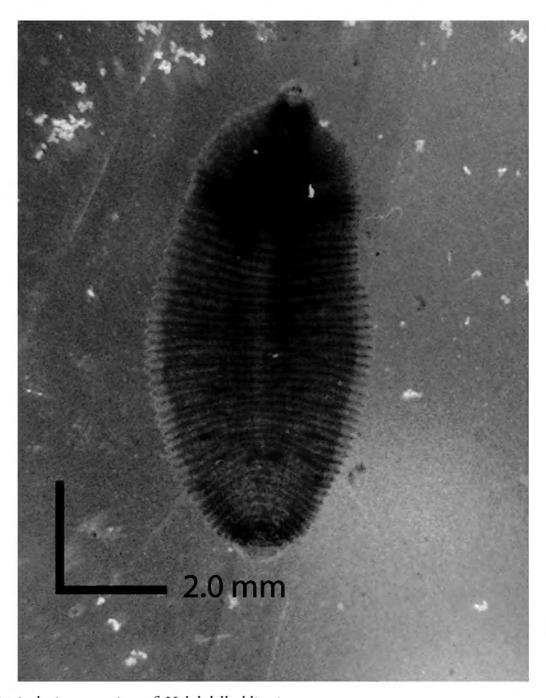


Figure 4. Typical pigmentation of *Helobdella blinni* sp. n.

Internal morphology. Average oral sucker diameter is 0.7 + 0.19 mm (N = 15), proboscis length is 3.5 + 1.1 mm (N = 17) (Table 3). Diffuse salivary glands are located near the anterior of the first pair of crop caecae. There are five pairs of smooth crop caecae and one lobed pair of posteriorly directed post caecae. Six pairs of compact testisacs are located in between each of the crop caecae. The intestine contains four pairs of caecae, with the first two pairs anteriorly directed and the other two pairs posteriorly directed. The intestine leads into an unraised anus located two annuli from the caudal sucker (Figure 3).

Development and growth. This species breeds year-round with peaks in spring and fall. Our specimens had an average of 7 to 11 white eggs (diameter 0.5 + 0.15 mm, N = 7) fixed to their ventral surface. Laboratory collections (2007–10) of H. blinni documented the eggs hatching 1 to 2 weeks after ovipositing (Beresic-Perrins 2010). Once hatched, the young attach to the ventral surface of the parent, allowing the parent to hunt for food and feed the young, occasionally feeding along with them. Prey consists of oligochaetes and other invertebrates. The average number of young per

adult is 7 + 3.3 (N = 97) ranging from 1 to 14 offspring. The young remain attached to the parent for an additional four to five weeks after hatching. Once the juveniles leave the parent, they tend to aggregate together on rocks (Beresic-Perrins 2010).

Molecular analysis

A Bayesian inference phylogenetic tree of the COI sequence data is presented in Figure 5. We include the posterior probabilities and maximum-likelihood branch supports >50. The Arizona populations of *Helobdella* c.f. *modesta* formed a sister clade to *Helobdella* blinni sp. n., supported by both the Bayesian and parsimony analyses. The results of the uncorrected p-distance analysis revealed a difference of 13.3% (233 nucleotides included) between the two groups (Table 6). The two groups form a larger clade with *H. modesta* (Ohio), *H. stagnalis* (UK), and *H. modesta* (Washington) which is supported by both Bayesian inference and maximum-likelihood. *H. blinni* differed from *H. modesta* (Ohio) by 13.7%, *H. stagnalis* (UK) by 16.3%, and *H. modesta* (Washington) by 16.3% (Table 6).

When we aligned all 78 sequences, there were four, ten-codon deletions within all of the Arizona sequences and *H. atli* (Oceguera-Figueroa and León-Regagnon 2005, Oceguera-Figueroa et al. 2010). When we performed the parsimony analysis, we included deletions as a 5th state in our first analysis and in our second, we treated the deletions as missing data. In the resulting 5th state tree, the two Arizona species remained sister taxa (100% support), but included in the clade was *H. atli* (100% and 58% support). The missing data tree placed *H. blinni* ancestral to *H. modesta* (Washington), *H. modesta* (Ohio), *H. stagnalis* (UK), and *H. c.f. modesta* with 100% branch support (Fig. 5).

Discussion

Helobdella blinni sp. n. has morphological and life-history traits similar to other Helobdella species, including possession of a nuchal scute, diffuse salivary glands, six pairs of testisacs, and extended parental care for the young (6–7 weeks; Tables 4, 5). Helobdella blinni, H. bowermani (Moser et al. 2013), H. octatestisaca (Lai et al. 2009), and H. c.f. modesta each have five pairs of smooth crop caecae as opposed to six pairs of lobed crop caecae in H. californica (Kutschera 2011) and H. papillornata (Govedich and Davies 1998). Helobdella blinni and H. temiscoensis (Salas-Montiel et al. 2014) share pigmentation characteristics, but they differ internally. Helobdella temiscoensis has only four pairs of crop caecae and one descending post caecae as opposed to five pairs and one descending post caecae in H. blinni. Helobdella modesta, H. californica, H. atli, H. bowermani, and H. octatestisaca do not resemble the pigmentation of H. blinni, running the spectrum from grey to pink. Additionally, they have a descending pair of post caecae, whereas H. atli, H. californica, and H. papillornata do not. Helobdella blinni,

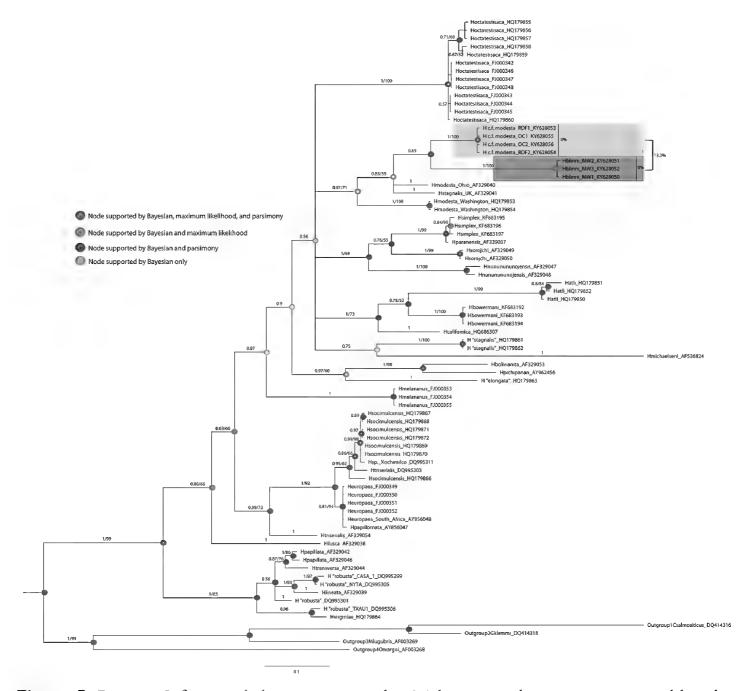


Figure 5. Bayesian Inference phylogenetic tree with 25% burn-in and support was assessed based on clade posterior probabilities tree. We included COI sequences from 31 species of *Helobdella* (family Glossiphoniidae). The Arizona populations are from Oak Creek (OC), Rio de Flag (RDF), and Montezuma Well (MW). Our outgroup included *Cystobranchus salmositicus* (Meyer, 1946), *Gonimosobdella klemmi* (Williams & Burreson, 2005), *Myzobdella lugubris* (Leidy, 1851), and *Ozobranchus margoi* (Davies, 1978). The shaded branches are the Arizona sample sequences. Branch labels include the Bayesian / ML probability. The blue nodes are supported by Bayesian Inference, Maximum-Likelihood, and parsimony analyses. The yellow nodes are supported by Bayesian Inference and Maximum-Likelihood analyses. The green nodes are supported by Bayesian Inference and parsimony analyses. The red nodes are supported by Bayesian Inference analysis only.

H. c.f. modesta, H. californica, H. temiscoensis, H. atli, and H. bowermani possess six pairs of testisacs, whereas H. papillornata has five pairs and H. octatestisaca has four pairs. Helobdella blinni also has a larger proboscis than the other Helobdella species (mean + SE, H. blinni 3.5 mm + 1.1, N=17, H. californica mean = 0.7 mm, H. papillornata mean= 2 mm). Furthermore, breeding periods also differ between H. blinni and the other Helobdella species (Tables 4, 5).

Table 4. Morphological comparison of Helobdella species

Traits	H. blinni sp. n.	H. modesta	H. californica	H. papillornata	H. temiscoensis	H. atli	H. bowermani	H. octatestisaca
	(current paper)	(Kutschera 1988: Sawyer 1986)	(Kutschera 1988; 2011)	(Govedich and Davies 1998)	(Salas-Montiel et al. 2014)	(Oceguera-Figueroa and León-Regagnon 2005)	(Moser et al. 2013)	(Lai et al. 2009)
crop caecae	5 pairs, smooth	5 pairs, smooth	6 pairs, lobed	5–6 pairs, lobed	4 pairs	6 pairs	5 pairs, smooth	5 pairs
post caecae	1 pair	1 pair	none	none	1 pair	none	1 pair	1 pair
eyes	1 pair	1 pair	1 pair	1 pair)	1 pair	1 pair	1 pair	1 pair
distance between eyes	0.1 mm	۸٠	د.	0.06 mm	٠.	۸.	۸.	۸.
nuchal scute	yes	yes	yes	ou	yes	yes	yes	yes
pairs of testisacs	6 pairs	6 pairs	6 pairs	5 pairs	6 pairs	6 pairs	6 pairs	4 pair
salivary glands	diffuse	diffuse	۲.	diffuse	diffuse	٠.	diffuse	diffuse
proboscis length	3.5mm	٠.	0.7mm	2mm	۲.	۲.	۸.	۸.
color	transparent with spots to dark brown	transparent to light grey	dark grey	transparent with stripes and papillae	pale brown, blackish - on posterior and mid-body	white-yellowish	pale yellow/ buff, papillae present	brown, pale, gray, and pink
body length	11–22 mm	8–12 mm	10–18 mm	15–40 mm	7.9–13.6 mm	7.5 mm	5.2–9.7 mm	9–14 mm
feeding	small invertebrates	small invertebrates	small invertebrates	small invertebrates	۲.	٠.	۸.	small invertebrates
brooding period	6–7 weeks	6–7 weeks	3-4 weeks	4–6 weeks	۲.	٠.	۸.	۸.
egg color	white	pink	pink	pink	÷	¿	۲:	۸.
egg diameter	0.5 mm	۲٠	0.5 mm	0.2 mm	۲.	۲٠	۸.	۸.
# eggs	7–14	12–35	8–56	20–50	۸.	۸.	۸.	۸.
1								

Table 5. Differences in brooding season and size between *H. blinni* sp. n., *H. stagnalis*, and *H.* c.f. *modesta*.

Location	Brooding Season	Average # of offspring	Author
H. blinni sp. n. Montezuma Well, AZ	Year-round	1–14	Beresic-Perrins (2010)
H. modesta Utah Lake, UT	Late spring through summer	12.6–17.4	Tillman and Barnes (1973)
H. modesta Lake Washington, WA	Spring and Summer	14.5	Thut (1969)
H. modesta Marion Lake, BC, CA	Spring and Summer	17.2–19.7	Davies and Reynoldson
<i>H. modesta</i> Newsome Pond, AB, CA	Late spring through summer	21.3	(1976)
<i>H. modesta</i> Cambridge, MA	Spring	31	Castle (1900)
H. modesta Michigan	Late spring through summer	35.3	Sawyer (1972)
H. stagnalis Iceland	Late spring through summer	No data	Bruun (1938)
H. stagnalis River Ely, South Wales	Late spring through summer	No data	Murphy and Learner (1982)
<i>H. stagnalis</i> Whiteknights Lake, UK	Late spring through summer	13–17	Mann (1957)
<i>H. stagnalis</i> Eglwys Nunydd, UK	Late spring through summer	14	Learner and Potter (1974)
<i>H. stagnalis</i> Denmark	Late spring through summer	20	Bennike (1943)

Helobdella blinni, unlike the other Helobdella species discussed here, breeds yearround, living in the thermally stable environment of Montezuma Well, with constant (19–24°C) year-round temperatures (Table 5). Monthly samples have individuals carrying cocoons every month of the year, with peak seasons in the spring and fall, a situation quite different than that for other Helobdella species, which have seasonally-constrained reproductive cycles, with egg-laying and brooding beginning in the spring and ending in the fall every year (Table 5). In addition to breeding year-round, H. blinni produces smaller broods (7–14 young) when compared to H. modesta and H. stagnalis (12–35 young) (Tables 4, 5) and has white eggs, unlike the characteristically pink eggs of H. modesta, H. californica, and H. papillornata (Table 4). The external pigmentation of H. blinni also tends to be dark brown, whereas most other Helobdella species are grey/brown in color (Fig. 4). Helobdella blinni are slightly longer (body length 11–22 mm, 16.6 + 3.2, N=24) than H. c.f. modesta (8–12 mm) and H. californica (10–18 mm), but slightly shorter in length than H. papillornata (15–40 mm) (Table 4).

The results from our molecular analysis show *H. blinni* to be genetically distinct from other *Helobdella* species, both from the same region (Rio de Flag and Oak Creek,

Table 6. Uncorrected p-distance pairwise analysis.

Species	Distance - H. blinni	Distance - H. c.f. modesta
H. atli	14.1–15%	16.7%
H. bolivianita	18.5%	19.3%
H. bowermani	15.9%	15.5%
H. blinni	0.0%	13.3%
H. californica	16.7%	17.6%
H. c.f. modesta	13.3%	0.0%
H. elongata	18.0%	19.3%
H. europaea	15.5%	16.3%
H. fusca	19.3%	20.6%
H. lineata	16.3%	14.6%
H. melananus	16.3%	17.2%
H. michaelseni	23.2%	20.6%
H. modesta OH	13.7%	8.6%
H. modesta WA	16.3%	14.6%
H. nununununojensis	17.5–19%	17.5–19.7%
H. octatestisaca	19.7%	16.3%
H. papillata	17.6%	15.8–16.3%
H. papillornata	15.9%	16.7%
H. paranensis	16.3%	13.7%
H. pichipanan	17.2%	19.3%
H. robusta	17.6%	14.6%
H. aff robusta CASA	18.9%	17.2%
H. aff robusta NYTA	17.6%	15.9%
H. aff robusta TXAU1	17.2%	18.0%
H. simplex	16.3%	13.7–14.2%
H. socimulcensis	15.9%	16.7–17.2%
H. sorojchi	18–18.5%	17.6%
H. sp. Xochimilco	15.5%	17.2%
H. stagnalis	20.6%	17.2%
H. stagnalis UK	16.3%	11.6%
H. transversa	16.3%	15.0%
H. triserialis	15.9%	16.7–18.5%
H. virginiae	16.3%	16.7%
Outgroup1 C. salmositicus	24.9%	23.2%
Outgroup2 G. klemmi	21.5%	21.5%
Outgroup3 M. lugubris	18.0%	19.3%
Outgroup4 O. margoi	23.2%	20.2%

Arizona populations) and from Europe (UK sample). The sequences yielded a 13.3%–17.4% genetic difference between *H. blinni* and both the Arizona *H.* c.f. *modesta*, *H. modesta* (Ohio and Washington), and the United Kingdom *H. stagnalis* populations (Table 6). The three Arizona populations belong to their own separate clade, but are closely related to *H. stagnalis* (UK) and the *H. modesta*. *Helobdella atli*, *H. bowermani*,

and *H. californica* are located on separate branch tips, but they comprise what Oceguera-Figueroa et al. (2010) designated as the "*stagnalis*" series (Fig. 5).

Based on morphological, life-history, and molecular differences, we propose the *Helobdella* sp. leeches found at Montezuma Well should be considered a new species, likely the result of allopatric isolation. This concept supports our hypothesis that the leech species inhabiting Montezuma Well may have become isolated from other populations as far back as 11,000 years ago (Wagner and Blinn 1987). *Helobdella blinni* sp. n. can be considered a distinct species found in Montezuma Well *and* may also turn out to be endemic to the area. Further sampling and analyses are needed in order to verify endemism.

Although currently classified as *Helobdella* c.f. *modesta*, the Arizona populations from the Rio de Flag and Oak Creek may be an additional undescribed species based on our molecular analysis. Our next step is to investigate these populations more closely, comparing them to other local populations, including White Horse Lake and J.D. Dam Lake, AZ which also contain *H. modesta*.

Acknowledgements

We would like to thank the National Park Service rangers and volunteers at the Montezuma Castle National Monument, especially Rex Vanderford, who assisted with collections and permits. Funding for the molecular and morphological analyses was provided by the National Science Foundation's Integrative Graduate Education and Research Traineeship. William Moser, Smithsonian Institution, National Museum of Natural History, Department of Invertebrate Zoology, provided valuable assistance in specimen management. Dr. Tina Ayers and Tamara Max (Northern Arizona University, Department of Biological Sciences) provided valuable assistance and insight with molecular data analyses. All the members past and present of Dr. Stephen Shuster's laboratory were indispensable with all aspects of data collection for this manuscript.

References

- Apakupakul K, Siddall ME, Burreson EM (1999) Higher level relationships of leeches (Annelida: Clitellata: Euhirudinea) based on morphology and gene sequences. Molecular Phylogenetics and Evolution 12: 350–359. https://doi.org/10.1006/mpev.1999.0639
- Bennike SAB (1943) Contributions to the ecology and biology of the Danish freshwater leeches (Hirudinea). Folia Limnologica Scandinavica 2: 1–109.
- Beresic-Perrins RK (2010) The life-history and parental care in an Arizona population of *Helob-della stagnalis* (Hirudinea: Glossiphoniidae). Master's thesis, Flagstaff, Arizona: Northern Arizona University.
- Bely AE, Weisblat DA (2006) Lessons from leeches: a call for DNA barcoding in the lab. Evolution & Development 8: 491–501. https://doi.org/10.1111/j.1525-142X.2006.00122.x

- Blanchard R (1896) Viaggio del Dott. A. Borellinella Republica Argentina e nel Paraguay. 21. Hirudine'es. Bollettino dei Musei di Zoologia ed. Anatomia comparata Universita di Torino 11: 1–24.
- Bruun AF (1938) Freshwater Hirudinea. Zoology Iceland 2: 22.
- Castle RE (1900) Some North American fresh-water Rhynchobdellida, and their parasites. Bulletin of the Museum of Comparative Zoology 36: 39–42.
- Cole GA, Barry WT (1973) Montezuma Well, Arizona, as a habitat. Journal of the Arizona Academy of Sciences 8: 7–13. https://doi.org/10.2307/40022206
- Cole GA, Watkins RL (1977) *Hyalella montezuma*, a new species (Crustacea: Amphipoda) from Montezuma Well, Arizona. Hydrobiologia 52:175–184. https://doi.org/10.1007/BF00036441
- Davies RW, Reynoldson TB (1976) A comparison of the life cycle of *Helobdella stagnalis* (Linn. 1758) (Hirudinoidea) in two different geographical areas. Canadian Journal of Animal Ecology 45: 457–470. https://doi.org/10.2307/3885
- Davies R (1978) The morphology of *Ozobranchus margoi* (Apathy) (Hirudinoidea), a parasite of marine turtles. The Journal of Parasitology 64: 1092–1096. https://doi.org/10.2307/3279733
- Davies RW, Singhal RN, Blinn DW (1985) *Erpobdella montezuma* (Hirudinoidea: Erpobdellidae), a new species of freshwater leech from North America. Canadian Journal of Zoology 63: 965–969. https://doi.org/10.1139/z85-142
- Edgar RC (2004) MUSCLE: multiple sequence alignment with high accuracy and high throughput. Nucleic Acids Research 32: 1792–1797. https://doi.org/10.1093/nar/gkh340
- Folmer O, Black M, Hoeh W, Lutz R, Vrijenhoek R (1994) DNA primers for amplification of mitochondrial cytochrome *C* oxidase subunit I from diverse metazoan invertebrates. Molecular Marine Biology and Biotechnology 3: 294–299.
- Govedich FR, Blinn DW, Keim P, Davies RW (1998) Phylogenetic relationships of three genera of Erpobdellidae (Hirudinoidea), with a description of a new genus, *Motobdella*, and species, *Motobdella sedonensis*. Canadian Journal of Zoology 76: 2164–2171. https://doi.org/10.1139/z98-132
- Govedich FR, Davies RW (1998) The first record of the genus *Helobdella* (Hirudinoidea: Glossiphoniidae) from Australia, with a description of a new species, *Helobdella papillornata*. Hydrobiologia 389: 45–49. https://doi.org/10.1023/A:1003543314841
- Kumar S, Stecher G, Tamura K (2016) MEGA7: Molecular Evolutionary Genetics Analysis version 7.0 for bigger datasets. Molecular Biology and Evolution 33: 1870–1874. https://doi.org/10.1093/molbev/msw054
- Kutschera U (1988) A new leech species from North America, *Helobdella californica* nov. sp. (Hirudinea: Glossiphoniidae). Zoologischer Anzeiger 220: 173–178.
- Kutschera U (2011) The Golden Gate leech *Helobdella californica* nov. sp. (Hirudinea: Glossiphoniidae): occurrence and DNA-based taxonomy of a species restricted to San Francisco. International review of Hydrobiology 96: 286–295. https://doi.org/10.1002/iroh.201111311
- Lai Y-T, Chang C-H, Chen J-H (2009) Two new species of *Helobdella* Blanchard 1896 (Hirudinida: Rhynchobdellida: Glossiphoniidae) from Taiwan, with a checklist of hirudinea fauna of the island. Zootaxa 2068: 27–46.

- Lanave C, Preparata G, Sacone C, Serio G (1984) A new method for calculating evolutionary substitution rates. Journal of Molecular Evolution 20: 86–93. https://doi.org/10.1007/BF02101990
- Lanfear R, Calcott B, Ho SY, Guindon S (2012) PartitionFinder: combined selection of partitioning schemes and substitution models for phylogenetic analyses. Molecular Biology and Evolution 29: 1695–701. https://doi.org/10.1093/molbev/mss020
- Learner MA, Potter DWB (1974) Life history and production of the leech *Helobdella stagnalis* (L.) (Hirudinea) in a shallow eutrophic reservoir in South Wales. Journal of Animal Ecology 43: 199–208. https://doi.org/10.2307/3167
- Leidy J (1851) Gen. nov. *Myzobdella*. Proceedings of the Academy of Natural Sciences of Philadelphia 5: 243.
- Linnaeus C (1758) Systema naturae per regna tria naturae, secundum classes, ordines, genera, species, cum characteribus, differentiis, synonymis, locis. Tomus I. Editio Decima. Holmiae, Laurentii Salvii.
- Mann KH (1957) The breeding, growth and age structure of a population of the leech *Helobdella stagnalis* (L.). Journal of Animal Ecology 26: 171–177. https://doi.org/10.2307/1787
- Meyer M (1946) Further Notes on the leeches (Piscicolidae) living on fresh-water fishes of North America. Transactions of the American Microscopical Society 65: 237–249. https://doi.org/10.2307/3223419
- Miller MA, Pfeiffer W, Schwartz T (2010) Creating the CIPRES Science Gateway for inference of large phylogenetic trees. In: Proceedings of the Gateway Computing Environments Workshop (GCE), New Orleans, LA, 1–8. https://doi.org/10.1109/gce.2010.5676129
- Moore JP (1898) Leeches of the United States National Museum. Proceedings of the U.S. National Museum 21: 543–563. https://doi.org/10.5479/si.00963801.21-1160.543
- Moore JP (1952) Professor A.E. Verrill's freshwater leeches a tribute and critique. Notulae Naturae of the Academy of Natural Sciences of Philadelphia 245: 1–15.
- Moser WE, Klemm DJ, Richardson DJ, Wheeler BA, Trauth SE, Daniels BA (2006) Leeches (Annelida: Hirudinida) of northern Arkansas. Journal of the Arkansas Academy of Science 60: 84–95.
- Moser WE, Richardson DJ, Hammond CI, Lazo-Wasem E (2011) Molecular characterization of *Helobdella modesta* (Verrill, 1872) (Hirudinida: Glossiphoniidae) from its type locality, West River and Whitneyville Lake, New Haven County, Connecticut, USA. Zootaxa 2834: 65–68.
- Moser WE, Fend SV, Richardson DJ, Hammond CI, Lazo-Wasem EA, Govedich FR, Gullo BS (2013) A new species of *Helobdella* (Hirudinida: Glossiphoniidae) from Oregon, USA. Zootaxa 3718: 287–94. https://doi.org/10.11646/zootaxa.3718.3.5
- Murphy PM, Learner MA (1982) The life history and reproduction of the leech *Helobdella stagnalis* (Hirudinea: Glossiphonidae) in the River Ely, South Wales. Freshwater Biology 12: 321–329. https://doi.org/10.1111/j.1365-2427.1982.tb00626.x
- Oceguera-Figueroa A, León-Regagnon V (2005) A new freshwater leech species of *Helob-della* (Annelida: Glossiphoniidae) from central Mexico. Zootaxa 976: 1–8. https://doi.org/10.11646/zootaxa.976.1.1

- Oceguera-Figueroa A, León-Règagnon V, Siddall ME (2010) DNA barcoding reveals Mexican diversity within the freshwater leech genus *Helobdella* (Annelida: Glossiphoniidae). Mitochondrial DNA 21: 24–9. https://doi.org/10.3109/19401736.2010.527965
- Rodriguez FJ, Oliver JL, Marin A, Medina JR (1990) The general stochastic model of nucleotide substitution. Journal of Theoretical Biology 142: 485–501. https://doi.org/10.1016/S0022-5193(05)80104-3
- Ronquist F, Huelsenbeck JP (2003) MrBayes 3: Bayesian phylogenetic inference under mixed models. Bioinformatics 19: 1572–1574. https://doi.org/10.1093/bioinformatics/btg180
- Salas-Montiel R, Phillips AJ, De Leon GP, Oceguera-Figueroa A (2014) Description of a new leech species of *Helobdella* (Clitellata: Glossiphoniidae) from Mexico with a review of Mexican congeners and a taxonomic key. Zootaxa 3900: 77–94. https://doi.org/10.11646/zootaxa.3900.1.4
- Sawyer RT (1972) North American freshwater leeches, exclusive of the Piscicolidae, with a key to all species. Illinois Biological Monographs, Univ. of Illinois Press, Urbana. https://doi.org/10.5962/bhl.title.53881
- Sawyer RT (1986) Leech Biology and Behaviour, Vols. I–III. Clarendon Press, Oxford, United Kingdom, 1065 pp.
- Siddall ME, Burreson EM (1998) Phylogeny of leeches (Hirudinea) based on mitochondrial cytochrome *c* oxidase subunit I. Molecular Phylogenetics and Evolution 9: 156–62. https://doi.org/10.1006/mpev.1997.0455
- Siddall ME, Borda E (2002) Phylogeny and revision of the leech genus *Helobdella* (Glossiphoniidae) based on mitochondrial gene sequences and morphological data and a special consideration of the *triserialis* complex. Zoologica Scripta 32: 23–33. https://doi.org/10.1046/j.1463-6409.2003.00098.x
- Siddall ME, Budinoff RB (2005) DNA-barcoding evidence for widespread introductions of a leech from the South American *Helobdella triserialis* complex. Conservation Genetics 6: 467–72. https://doi.org/10.1007/s10592-005-4986-y
- Siddall ME, Budinoff RB, Borda E (2005) Phylogenetic evaluation of systematics and biogeography of the leech family Glossiphoniidae. Invertebrate Systematics 19: 105–112. https://doi.org/10.1071/IS04034
- Stamatakis A (2014) RAxML Version 8: A tool for phylogenetic analysis and post-analysis of large phylogenies. Bioinformatics 30: 1312–1313. https://doi.org/10.1093/bioinformatics/btu033
- Swofford DL (2003) PAUP* Version 4. Phylogenetic analysis using parsimony (*and other methods).
- Tavare S (1986) Some probabilistic and statistical problems on the analysis of DNA sequences. In: Lectures in mathematics in the life sciences, vol. 17, 57–86.
- Thut RN (1969) A study of the profundal bottom fauna of Lake Washington. Ecological Monographs 39: 79–100. https://doi.org/10.2307/1948566
- Tillman DL, Barnes JR (1973) The reproductive biology of the leech *Helobdella stagnalis* (L.) in Utah Lake, Utah. Freshwater Biology 3: 137–145. https://doi.org/10.1111/j.1365-24-27.1973.tb00068.x

- Vaillant L (1890) Histoire naturelle des Anneles marin et d'eau douce. 3/2, Ordre Hirudiens (Hirudines) ou Bdelles, Paris, 477–542.
- Verrill AE (1872) Descriptions of North American freshwater leeches. American Journal of Science and Arts 3:126–139. https://doi.org/10.2475/ajs.s3-3.14.126
- Wagner VT, Blinn DW (1987) Montezuma Well: the living desert oasis. http://www.nps.gov/history/online_books/moca/montezuma_well/index.htm
- Williams JI, Burreson EM (2005) *Gonimosobdella klemmi* n. gen., n. sp. (Hirudinida: Piscicolidae) from cyprinid fishes in Arkansas, Illinois, and Missouri, USA. Comparative Parasitology 72: 166–72. https://doi.org/10.1654/4180
- Williams JI, Burreson EM (2006) Phylogeny of the fish leeches (Oligochaeta, Hirudinida, Piscicolidae) based on nuclear and mitochondrial genes and morphology. Zoologica Scripta 35: 627–639. https://doi.org/10.1111/j.1463-6409.2006.00246.x